## AMENDMENTS TO THE CLAIMS

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- 1. (Previously presented) A method of increasing the cutting hardness of a shaped body comprising a crystalline aluminosilicate having an SiO<sub>2</sub>/Al<sub>2</sub>O<sub>3</sub> molar ratio of 10:1 to 1200:1, wherein the shaped body comprises a binder selected from among oxides of silicon and/or zirconium and is treated with a gas consisting of water vapor at from 100 to 600°C and an absolute pressure of from 0.1 to 10 bar for a period of at least 20 hours and the shaped body has been calcined at from 100 to 600°C before the treatment with water vapor.
- 2. (Previously presented) The method according to claim 1, wherein the shaped body is treated for a period of at least 50 hours.
- 3. (Previously presented) The method according to claim 1, wherein the shaped body is treated continuously at a WHSV (weight hourly space velocity) of from 0.05 to 5 g of water vapor per gram of shaped body and per hour (g<sub>water vapor</sub>/(g<sub>shaped body</sub> h)).
- 4. (Previously presented) The method according to claim 1, wherein the shaped body is treated continuously at a WHSV (weight hourly space velocity) of from 0.1 to 1 g of water vapor per gram of shaped body and per hour (gwater vapor/(gshaped body h)).
- 5. (Previously presented) The method according to claim 1, wherein the shaped body is treated at from 200 to 450°C and an absolute pressure of from 0.1 to 2 bar.
- 6. (Previously presented) The method according to claim 1, wherein the shaped body is fixed in position (fixed bed) during the treatment with water vapor.
- 7. (Cancelled)
- 8. (Previously presented) The method according to claim 1, wherein the crystalline aluminosilicate in the shaped body has an SiO<sub>2</sub>/Al<sub>2</sub>O<sub>3</sub> molar ratio of greater than 50:1.
- 9. (Previously presented) The method according to claim 1, wherein the crystalline aluminosilicate in the shaped body is at least partly in the H<sup>+</sup> and/or NH<sub>4</sub><sup>+</sup> form.

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10. (Previously presented) The method according to claim 1, wherein the crystalline aluminosilicate in the shaped body is of the pentasil type.

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- 11. (Previously presented) A process for preparing triethylenediamine (TEDA) by reaction of ethylenediamine (EDA) and/or piperazine (PIP) in the presence of a crystalline aluminosilicate catalyst, wherein a shaped body whose cutting hardness has been increased beforehand using a method according to claim 1 is used as catalyst.
- 12. (Previously presented) The process according to claim 11, wherein the reaction is carried out continuously and in the gas phase.
- 13. (Previously presented) The process according to claim 11, wherein EDA and one or more amine compounds selected from the group consisting of monoethanolamine, diethanolamine, triethanolamine, PIP, diethylenetriamine, triethylenetetramine, tri(2-aminoethyl)amine, morpholine, N-(2-aminoethyl)ethanolamine, N-(2-hydroxyethyl)piperazine, N-(2-aminoethyl)piperazine, N,N'-bis(2-hydroxyethyl)piperazine and N-(2-aminoethyl)-N'-(2-hydroxyethyl)piperazine are reacted.
- 14. (Previously presented) The process according to claim 11, wherein EDA and from 7 to 250% by weight of piperazine (PIP), based on EDA, are reacted.
- 15. (Previously presented) The process according to claim 11, wherein EDA, from 8 to 250% by weight of PIP and from 23 to 300% by weight of water, in each case based on EDA, are reacted.
- 16. (Previously presented) The process according to claim 11, wherein the reaction temperature for the reaction to form TEDA is from 310 to 390°C.
- 17. (Previously presented) The process according to claim 11, wherein the absolute pressure in the reaction to form TEDA is from 0.1 to 10 bar.
- 18. (Cancelled)

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After Final Office Action of January 15, 2009

(Previously presented) A process for chemical synthesis carried out in the presence of a 19. crystalline aluminosilicate catalyst, wherein a shaped body whose cutting hardness has been increased beforehand using a method according to claim 1 is used as catalyst.

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(Previously presented) The process according to claim 19, wherein the synthesis is an 20. alkylation, disproportionation, acylation, isomerization, oligomerization, amination, alkoxylation, epoxidation, cyclization, hydroxylation, condensation, hydration or dehydration.

## 21. (Cancelled)

22. (Previously presented) A shaped body prepared by the method as claimed in claim 1.

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